This presentation should not be considered a final statement of NIOSH policy or of any agency or individual who was involved. This information is intended for use in advancing knowledge needed to protect workers. Comments regarding this presentation may be submitted to the NIOSH Docket Office.

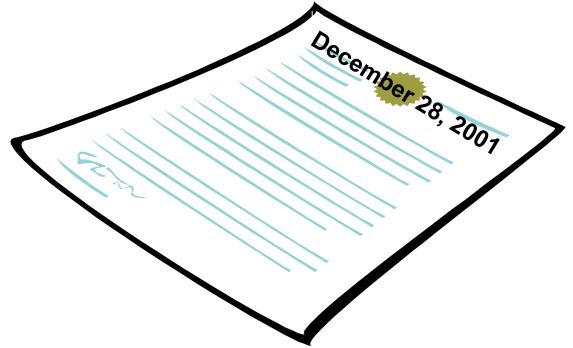


NIOSH Public Meeting

Proposed CBRN
Full- Facepiece
APR Standard



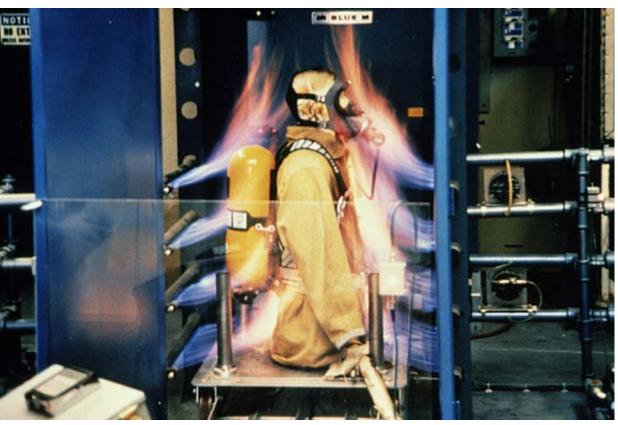
CBRN Standard for SCBA



NIOSH will follow a flexible program for test and evaluation of SCBA for a limited period of 1 year.



CBRN Standard for SCBA





CBRN Standard for SCBA

Unable to:

Test against Sarin and Mustard Gas

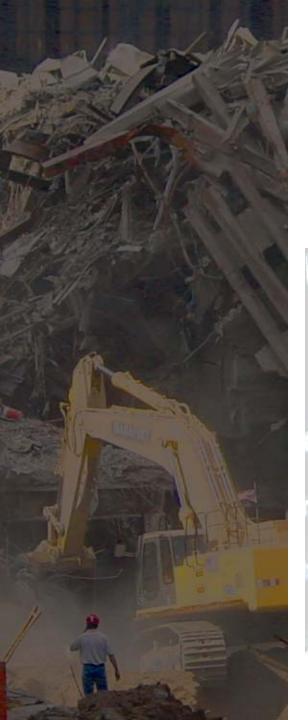
Reliably predict testing outcome

Sample products to meet requirements



CBRN Standard for SCBA





CBRN Standard for SCBA

Identify & Recognize Surrogate Agents

Continue flexible SCBA CBRN approval process.

Follow a similar path for the CBRN full facepiece air purifying respirator standard.



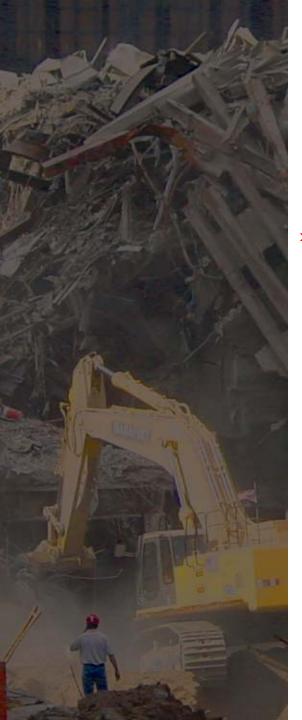
Publicly Validate Protocol

- Results and test repeatability must be trusted
- CBRN SCBA protocol was inadequately validated
- Thoroughly validate the process and publish the results



Opportunity to Comment

- Most likely use scenario and rationale behind conditions
- Avoid unacceptable market conditions



Identification and acceptance of surrogate agents





NIOSH should not regulate interchangeability

Section (6)(a)

Interchangeable consumable filter cartridges and canisters







Reflections on September 11th

- Transportation infrastructure in the United
 States was shut down
- PPE needs filled locally
- Improve availability by creating stockpiles
- 100,000 Respirators & 200,000 Cartridges shipped to WTC in first 48 hours
- Not a supply issue, logistic & training issue

Reflections on September 11th

- Multiple government agencies were taking responsibility
- Rescue workers did not know where to get the products
- Not familiar with product use and maintenance requirements
- Develop an emergency logistics program to address large-scale events



- NIOSH should not regulate interchangeability
- MSA fully supports the comments by the ISEA
- Respirator developed as a system
- Mixing manufacturer components will not provide the desired results



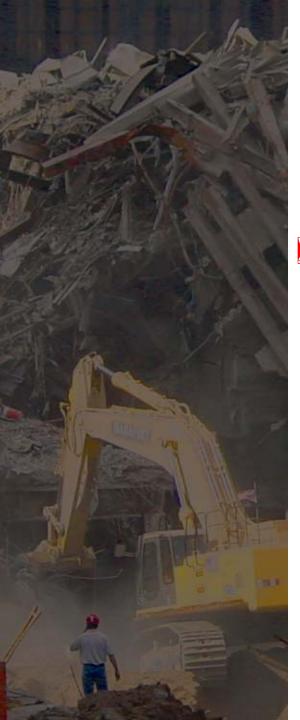
NIOSH should not regulate interchangeability





NIOSH should not regulate interchangeability

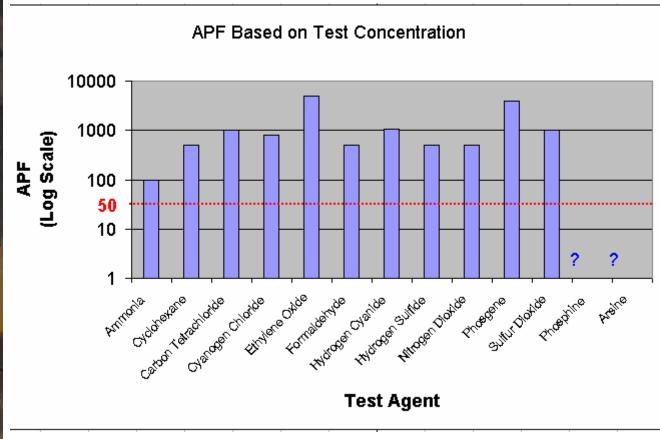
- NIOSH should maintain a systems level approval approach focusing on performance, not design
- NIOSH should support the development of emergency logistics programs



- NIOSH should not recommend or approve air-purifying respirators for entrance or use in IDLH or unknown atmospheres
- Beyond accepted use conditions of Air Purifying Respirators
- Forces test concentrations beyond accepted Assigned Protection Factors



CBRN APR Concerns





NIOSH should not recommend or approve air-purifying respirators for entrance or use in IDLH or unknown atmospheres

 SCBA with chemically protective clothing are the only responsible recommendation



NIOSH should follow a formal rule making process

- Details of the standard not final
- New respirator category worthy of a formal rule making process



- MSA supports the comments provided by the ISEA
- Recommend that NIOSH positions search for acceptable surrogates as their number one priority
- Maintain focus on performance requirements
- Do not approve any APR for entrance or use in IDLH or unknown atmospheres
- Follow formal rule making process



NIOSH Public Meeting

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